DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: S & B PHARMA, INC.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before [INSERT 60 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESS: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

The Attorney General has delegated his authority under the Controlled Substances

Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR

0.100(b). Authority to exercise all necessary functions with respect to the promulgation
and implementation of 21 CFR part 1301, incident to the registration of manufacturers,
distributors, and dispensers of controlled substances (other than final orders in connection
with suspension, denial, or revocation of registration) has been redelegated to the Deputy

Assistant Administrator of the DEA Office of Diversion Control ("Deputy Assistant

Administrator") pursuant to sec. 7(g) of 28 CFR pt. 0, subpt. R, App.

In accordance with 21 CFR 1301.33(a), this is notice that on December 10, 2013, S & B Pharma, Inc., DBA Norac Pharma, 405 South Motor Avenue, Azusa, California

91702-3232, applied to be registered as a bulk manufacturer of the following basic classes of narcotic or nonnarcotic controlled substances:

| Controlled Substance | Schedule |
|------------------------------------|-----------------|
| | |
| Gamma Hydroxybutyric (2010) | I |
| Tetrahydrocannabinols (7370) | I |
| Methamphetamine (1105) | II |
| Pentobarbital (2270) | II |
| Nabilone (7379) | II |
| 4-Anilino-N-phenethyl-4-piperidine | |
| (8333) | II |
| Tapentadol (9780) | II |
| Fentanyl (9801) | II |

The company plans to manufacture the listed controlled substances in bulk for use in product development and for commercial sales to its customers.

Dated: July 1, 2014.

Joseph T. Rannazzisi, Deputy Assistant Administrator.

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